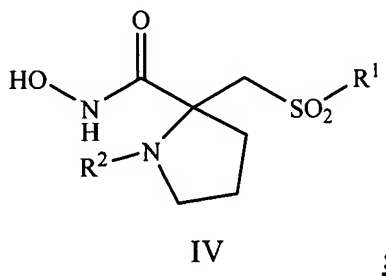


Amended Claims

Claims 1-14 (canceled)

15. (currently amended) A compound or a salt thereof, wherein:  
the compound corresponds in structure corresponding to Formula IV:



**wherein**

R<sup>2</sup> is selected from the group consisting of hydrido, C<sub>1</sub>-C<sub>8</sub> hydrocarbonyl, C<sub>1</sub>-C<sub>6</sub> hydrocarbyloxycarbonyl C<sub>1</sub>-C<sub>4</sub> hydrocarbonyl, aryl C<sub>1</sub>-C<sub>4</sub> hydrocarbonyl, heteroaryl C<sub>1</sub>-C<sub>4</sub> hydrocarbonyl, aryloxy C<sub>1</sub>-C<sub>4</sub> hydrocarbonyl, and [[or]] heteroaryloxy C<sub>1</sub>-C<sub>4</sub> hydrocarbonyl; and

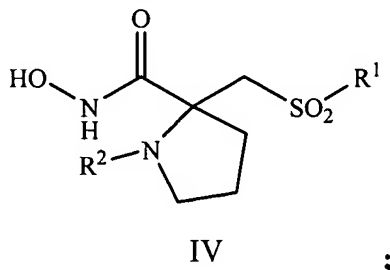
R<sup>1</sup> is ~~a substituent containing a phenyl single aryl~~ or 5- or 6-membered heteroaryl ~~radical~~ bonded directly to the depicted SO<sub>2</sub>-group that is ~~itself~~ substituted at its ~~own~~ 4-position when a 6-membered ring and at its ~~own~~ 3- or 4- position when a 5-membered ring with a substituent selected from the group consisting of ~~one other phenyl~~, single-ringed ~~aryl or~~ heteroaryl **group**, [[a]] C<sub>3</sub>-C<sub>14</sub> hydrocarbonyl **group**, [[a]] C<sub>2</sub>-C<sub>14</sub> hydrocarbyloxy **group**, [[a]] phenoxy **group**, [[a]] thiophenoxy **group**, [[a]] 4-thiopyridyl **group**, [[a]] phenylazo **group**, [[a]] phenylureido **group**, [[a]] nicotinamido **group**, [[an]] isonicotinamido **group**, [[a]] picolinamido **group**, [[an]] aniline, **group** and benzamido **group**.

16. (currently amended) The compound or salt according to claim 15, wherein:  
~~said R<sup>1</sup> radical is PhR<sup>3</sup> in which Ph~~ is phenyl substituted with R<sup>3</sup> at the 4-position; [[,]]  
and

R<sup>3</sup> is selected from the group consisting of [[a]] phenyl, phenoxy, thiophenoxy, anilino, phenylazo, benzamido, nicotinamido, isonicotinamido, picolinamido, and [[or]] phenylureido **group**.

17. (currently amended) ~~The A compound or a salt thereof according to claim 15,~~  
wherein:

the compound corresponds in structure to Formula IV:



R<sup>2</sup> is selected from the group consisting of hydrido, C<sub>1</sub>-C<sub>8</sub> hydrocarbyl, C<sub>1</sub>-C<sub>6</sub> hydrocarbyloxycarbonyl C<sub>1</sub>-C<sub>4</sub> hydrocarbyl, aryl C<sub>1</sub>-C<sub>4</sub> hydrocarbyl, heteroaryl C<sub>1</sub>-C<sub>4</sub> hydrocarbyl, aryloxy C<sub>1</sub>-C<sub>4</sub> hydrocarbyl, and heteroaryloxy C<sub>1</sub>-C<sub>4</sub> hydrocarbyl;

~~said R<sup>1</sup> radical is PhR<sup>3</sup> in which Ph is phenyl substituted with R<sup>3</sup> at the 4-position; [[,]]~~  
and

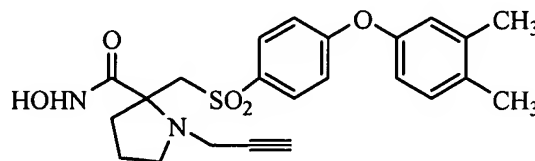
~~said R<sup>3</sup> is selected from the group consisting of [[a]] phenyl, phenoxy, anilino, thiophenoxy, benzamido, nicotinamido, isonicotinamido, picolinamido, and [[or]] phenylureido, wherein the phenyl, phenoxy, anilino, thiophenoxy, benzamido, nicotinamido, isonicotinamido, picolinamido, and phenylureido group that~~ is optionally substituted:

at the meta- or para-position or both with a ~~moiety that is~~ substituent selected from the group consisting of [[a]] halogen, [[a]] C<sub>1</sub>-C<sub>9</sub> hydrocarbyloxy **group**, [[a]] C<sub>1</sub>-C<sub>10</sub> hydrocarbyl **group**, [[a]] di-C<sub>1</sub>-C<sub>9</sub> hydrocarbylamino **group**, [[a]] carboxyl C<sub>1</sub>-C<sub>8</sub> hydrocarbyl **group**, [[a]] C<sub>1</sub>-C<sub>4</sub> hydrocarbyloxy carbonyl C<sub>1</sub>-C<sub>4</sub> hydrocarbyl **group**, [[a]] C<sub>1</sub>-C<sub>4</sub> hydrocarbyloxycarbonyl C<sub>1</sub>-C<sub>4</sub> hydrocarbyl, **group** and [[a]] carboxamido C<sub>1</sub>-C<sub>8</sub> hydrocarbyl **group**, or ~~is substituted~~

at the meta- and para-positions by two methyl groups or by methylenedioxy **group**.

Claims 18-24 (canceled)

25. (currently amended) A compound or a salt thereof, wherein the compound corresponds ~~corresponding~~ in structure to the formula:



**Claims 26-31 (canceled)**

32. (new) A method for treating a mammal having a condition associated with matrix metalloproteinase activity, wherein the method comprises administering to the mammal a therapeutically-effective amount of a compound recited in claim 15 or a pharmaceutically-acceptable salt thereof.

33. (new) A method for treating a mammal having a condition associated with matrix metalloproteinase activity, wherein the method comprises administering to the mammal a therapeutically-effective amount of a compound recited in claim 17 or a pharmaceutically-acceptable salt thereof.

34. (new) A method for treating a mammal having a condition associated with matrix metalloproteinase activity, wherein the method comprises administering to the mammal a therapeutically-effective amount of a compound recited in claim 25 or a pharmaceutically-acceptable salt thereof.